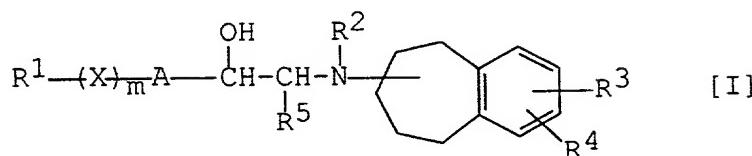


## CLAIMS

1. A compound of the general formula [I] :

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wherein

10       $R^1$  is aryl which may have one or more suitable  
          substituent(s), heterocyclic group or  
          cyclo(lower)alkyl,

15       $R^2$  is hydrogen or amino protective group,

20       $R^3$  and  $R^4$  are independently hydrogen, halogen, hydroxy,  
          amino, nitro, carboxy, protected carboxy, aryl,  
          lower alkyl, hydroxy(lower)alkyl,  
          amino(lower)alkyl, acyloxy(lower)alkyl,  
          acylamino(lower)alkyl, lower alkylamino(lower)alkyl  
          which may have one or more suitable substituent(s),  
          mono or di-(lower)alkylamino, acylamino, acyl  
          group, lower alkoxy, halo(lower)alcoxy, lower  
          alkenyloxy, lower alkoxy(lower)alcoxy, aryloxy,  
          cyclo(lower)alkyloxy, heterocyclicoxy,  
          ar(lower)alkyloxy, acyloxy or acyl(lower)alcoxy,

25       $R^5$  is hydrogen, lower alkyl, or aryl,

A is lower alkylene which may have one or more suitable  
          substituent(s) or lower alkenylene,

X is O, S, SO, SO<sub>2</sub> or NH, and

m is an integer of 0 or 1,

30      or a salt thereof.

2. A compound of claim 1, wherein

35       $R^1$  is phenyl which may have one or more suitable  
          substituent(s),

$R^2$  is hydrogen,

R<sup>3</sup> is acyl(lower)alkoxy, lower alkoxy, protected carboxy, hydroxy or acyloxy,

R<sup>4</sup> is hydrogen,

R<sup>5</sup> is hydrogen,

5 A is lower alkylene,

X is O, and

m is an integer of 1.

3. A compound of claim 2, wherein

10 R<sup>1</sup> is phenyl which may have 1 or 2 suitable substituent(s) selected from the group consisting of hydroxy and lower alkylsulfonylamino,

R<sup>3</sup> is lower alkylcarbamoyl(lower)alkoxy,

heterocycliccarbamoyl(lower)alkoxy,

15 N-lower alkyl-lower alkylcarbamoyl(lower)alkoxy,

hydroxy,

lower alkoxy,

protected carboxy,

20 arylcarbamoyl(lower)alkoxy which may have lower

alkoxy or di(lower)alkylamino,

di-lower alkylsulfamoyloxy,

N-lower alkyl-heterocyclic(lower)alkylcarbamoyl-(lower)alkoxy,

25 N-lower alkyl-lower alkylcarbamoyl(lower)alkoxy or

N-lower alkyl-cyclo(lower)alkylcarbamoyl(lower)-alkoxy.

4. A compound of claim 3, wherein

30 R<sup>1</sup> is phenyl which may have hydroxy and methylsulfonylamino,

R<sup>3</sup> is ethylcarbamoylmethoxy,

indolylcarbamoylmethoxy,

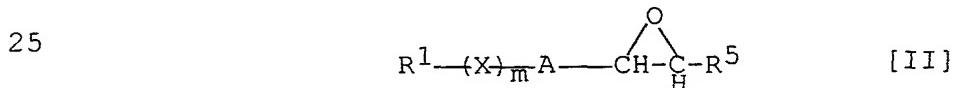
piperidinocarbonylmethoxy,

35 N-methylbutylcarbamoylmethoxy,

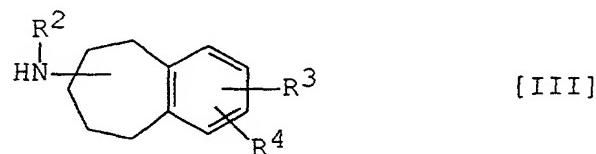
hydroxy,  
 butylcarbamoylmethoxy,  
 methoxy,  
 methoxycarbonyl,  
 5 ethoxy,  
 dimethylsulfamoyloxy,  
 tetrazolylcarbamoylmethoxy,  
 N-methylpyridylethylcarbamoylmethoxy,  
 methoxyphenylcarbamoylmethoxy,  
 10 thiazolylcarbamoylmethoxy,  
 dihydroindolylcarbonylmethoxy,  
 N-ethylpropylcarbamoylmethoxy,  
 N-methylbutylcarbamoylmethoxy,  
 N-ethylbutylcarbamoylmethoxy,  
 15 dimethylaminophenylcarbamoylmethoxy or  
 N-methylcyclohexylcarbamoylmethoxy.

5. A process for preparing a compound of claim 1,  
 or a salt thereof,  
 20 which comprises,

(i) reacting a compound [III] of the formula :

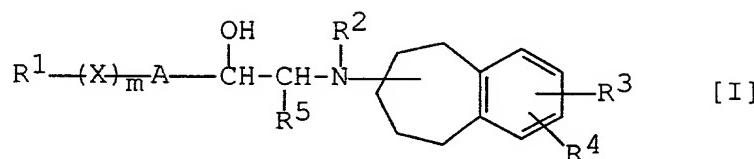


wherein  $\text{R}^1$ ,  $\text{R}^5$ ,  $\text{A}$ ,  $\text{X}$  and  $m$  are each as defined in  
 30 claim 1, with a compound [III] of the formula :



wherein R<sup>2</sup>, R<sup>3</sup> and R<sup>4</sup> are each as defined in claim 1,  
or a salt thereof, to give a compound [I] of the  
formula :

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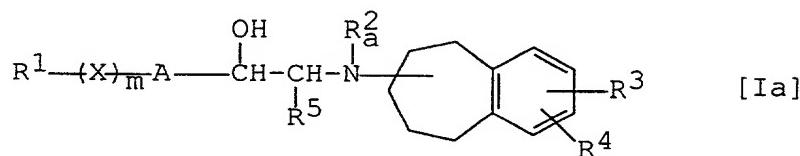
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wherein R<sup>1</sup>, R<sup>2</sup>, R<sup>3</sup>, R<sup>4</sup>, R<sup>5</sup>, A, X and m are each as  
defined in claim 1,  
or a salt thereof, or

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(ii) subjecting a compound [Ia] of the formula :

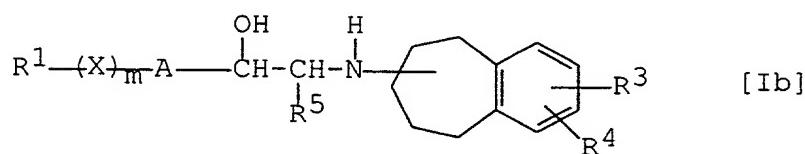
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wherein R<sup>1</sup>, R<sup>3</sup>, R<sup>4</sup>, R<sup>5</sup>, A, X and m are each as  
defined in claim 1, and

R<sup>2</sup><sub>a</sub> is amino protective group, or a salt thereof,  
to elimination reaction of the amino protective group,  
to give a compound [Ib] of the formula :

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wherein R<sup>1</sup>, R<sup>3</sup>, R<sup>4</sup>, R<sup>5</sup>, A, X and m are each as  
defined in claim 1,  
or a salt thereof.

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6. A pharmaceutical composition which comprises, as an  
active ingredient, a compound of claim 1 or a

pharmaceutically acceptable salt thereof in admixture with pharmaceutically acceptable carriers or excipients.

7. Use of a compound of claim 1 or a pharmaceutically acceptable salt thereof for the manufacture of a medicament.
8. A compound of claim 1 or a pharmaceutically acceptable salt thereof for use as a medicament.
9. A method for the prophylactic and/or the therapeutic treatment of pollakiuria or urinary incontinence which comprises administering a compound of claim 1 or a pharmaceutically acceptable salt thereof to a human being or an animal.

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